	U	1	Document ID	Issue Date	Pages	Title
1	×		US 20040067542 A1	20040408	18	P2y purinergic receptor expression for identifying preneoplastic and neoplastic states
2	Ø		US 20040018533 A1	20040129	107	Diagnosing predisposition to fat deposition and therapeutic methods for reducing fat deposition and treatment of associated
3	Ø		US 6242216 B1	20010605	40	Nucleic acids encoding a functional human purinoreceptor P2X2 and P2X4, and methods of production and use thereof
4	Ø		JP 2003093099 A	20030402 15 OSTEOCLAST DIFFERENTIATION-ASSOCIATE		OSTEOCLAST DIFFERENTIATION-ASSOCIATED GENE
5			WO 9741222 A1	19971106	32	HUMAN P2x4 RECEPTOR SPLICE-VARIANTS
6	×		US 6242216 B	20010605	40	Isolated polynucleotides, used to produce PX2 receptor polypeptides and identify potentially therapeutic compounds, encode a human P2X2 receptor polypeptide

	Current OR	Current XRef	Retrieval Classif	Inventor	s	С	P	2	3
1	435/7.23			Barden, Julian et al.					
2	435/6			Adam, Gail Isabel Reid et al.					
3	435/69.1	435/252.3; 435/320.1; 536/23.5		Lynch, Kevin J. et al.					
4				TAKEYA, TATSUO					
5				MCHALE, MARK THOMAS et al.	⊠				
6				BURGARD, E C et al.					

	4	5	Image Doc. Displayed	PT
1			US 20040067542	
2			US 20040018533	
3			US 6242216	
4			JP 2003093099 A	
5			WO 9741222 A1	
6			US 6242216	

```
Trying 31060000009999...Open
DIALOG INFORMATION SERVICES
PLEASE LOGON:
 ****** HHHHHHHH SSSSSSSS? ### Status: Signing onto Dialog *******
ENTER PASSWORD:
 ****** HHHHHHHH SSSSSSS? ******
Welcome to DIALOG
### Status: Login successfulDialog level 04.09.00D
Last logoff: 21may04 13:46:08
Logon file405 27may04 13:01:24
           *** ANNOUNCEMENT ***
--File 654 - US published applications from March 15, 2001 to the
present are now online. Please see HELP NEWS 654 for details.
--File 581 - The 2003 annual reload of Population Demographics is
complete. Please see Help News581 for details.
--File 990 - NewsRoom now contains February 2003 to current records.
File 992 - NewsRoom 2003 archive has been newly created and contains
records from January 2003. The oldest months's records roll out of
File 990 and into File 992 on the first weekend of each month.
To search all 2003 records BEGIN 990, 992, or B NEWS2003, a new
OneSearch category.
-- Connect Time joins DialUnits as pricing options on Dialog.
See HELP CONNECT for information.
                   ***
--SourceOne patents are now delivered to your email inbox
as PDF replacing TIFF delivery. See HELP SOURCE1 for more
information.
-- Important Notice to Freelance Authors--
See HELP FREELANCE for more information
NEW FILES RELEASED
***AeroBase (File 104)
***DIOGENES: Adverse Drug Events Database (File 181)
***World News Connection (File 985)
***Dialog NewsRoom - 2003 Archive (File 992)
***TRADEMARKSCAN-Czech Republic (File 680)
***TRADEMARKSCAN-Hungary (File 681)
***TRADEMARKSCAN-Poland (File 682)
UPDATING RESUMED
                    * * *
RELOADED
***Toxfile (File 156)
***Medline (Files 154-155)
***Population Demographics - (File 581)
***CLAIMS Citation (Files 220-222)
```

REMOVED

* * *

```
>>> Enter BEGIN HOMEBASE for Dialog Announcements <<<
>>> of new databases, price changes, etc.
```

* ALL NEW CURRENT YEAR RANGES HAVE BEEN * * *

SYSTEM: HOME

Cost is in DialUnits

Menu System II: D2 version 1.7.9 term=ASCII

*** DIALOG HOMEBASE(SM) Main Menu ***

Information:

- 1. Announcements (new files, reloads, etc.)
- 2. Database, Rates, & Command Descriptions
- 3. Help in Choosing Databases for Your Topic
- 4. Customer Services (telephone assistance, training, seminars, etc.)
- 5. Product Descriptions

Connections:

- 6. DIALOG(R) Document Delivery
- 7. Data Star(R)
 - (c) 2003 Dialog, a Thomson business.

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/H = Help

/L = Logoff

/NOMENU = Command Mode

Enter an option number to view information or to connect to an online service. Enter a BEGIN command plus a file number to search a database (e.g., B1 for ERIC).

Terminal set to DLINK

*** DIALOG HOMEBASE(SM) Main Menu ***

Information:

- 1. Announcements (new files, reloads, etc.)

- Database, Rates, & Command Descriptions
 Help in Choosing Databases for Your Topic
 Customer Services (telephone assistance, training, seminars, etc.)
 Product Descriptions

Connections:

- 6. DIALOG(R) Document Delivery
- 7. Data Star(R)
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/H = Help

/L = Logoff

/NOMENU = Command Mode

Enter an option number to view information or to connect to an online service. Enter a BEGIN command plus a file number to search a database (e.g., B1 for ERIC).

? b biochem, biotech

27may04 13:01:36 User276619 Session D5.1

0.155 DialUnits FileHomeBase

- \$0.00 Estimated cost FileHomeBase
- \$0.05 TELNET
- \$0.05 Estimated cost this search
- 0.155 DialUnits \$0.05 Estimated total session cost

```
SYSTEM:OS - DIALOG OneSearch
         5:Biosis Previews(R) 1969-2004/May W4
  File
         (c) 2004 BIOSIS
  File
         6:NTIS 1964-2004/May W4
        (c) 2004 NTIS, Intl Cpyrght All Rights Res 34:SciSearch(R) Cited Ref Sci 1990-2004/May W4
  File
         (c) 2004 Inst for Sci Info
        40:Enviroline(R) 1975-2004/Apr
  File
        50:CAB Abstracts 1972-2004/Apr
  File
         (c) 2004 CAB International
        65:Inside Conferences 1993-2004/May W4
  File
         (c) 2004 BLDSC all rts. reserv.
  File
        71:ELSEVIER BIOBASE 1994-2004/May W3
         (c) 2004 Elsevier Science B.V.
        73:EMBASE 1974-2004/May W4
  File
         (c) 2004 Elsevier Science B.V.
  File
        94:JICST-EPlus 1985-2004/May W1
         (c) 2004 Japan Science and Tech Corp(JST)
        98:General Sci Abs/Full-Text 1984-2004/May
         (c) 2004 The HW Wilson Co.
  File 103: Energy SciTec 1974-2004/May B1
         (c) 2004 Contains copyrighted material
*File 103: For access restrictions see Help Restrict.
  File 143:Biol. & Agric. Index 1983-2004/Apr
         (c) 2004 The HW Wilson Co
  File 144: Pascal 1973-2004/May W3
         (c) 2004 INIST/CNRS
  File 155:MEDLINE(R) 1966-2004/May W4
         (c) format only 2004 The Dialog Corp.
*File 155: Medline has been reloaded. Accession numbers
have changed. Please see HELP NEWS 154 for details.
  File 156:ToxFile 1965-2004/May W2
         (c) format only 2004 The Dialog Corporation
*File 156: ToxFile now reloaded with 2004 MeSH.
Enter Help News156 for more information.
  File 162:Global Health 1983-2004/Apr
         (c) 2004 CAB International
  File 172:EMBASE Alert 2004/May W3
         (c) 2004 Elsevier Science B.V.
  File 305: Analytical Abstracts 1980-2004/May W3
         (c) 2004 Royal Soc Chemistry
*File 305: Alert feature enhanced for multiple files, duplicate
removal, customized scheduling. See HELP ALERT.
  File 369: New Scientist 1994-2004/May W3
         (c) 2004 Reed Business Information Ltd.
  File 370:Science 1996-1999/Jul W3
         (c) 1999 AAAS
*File 370: This file is closed (no updates). Use File 47 for more current
information.
  File 399:CA SEARCH(R) 1967-2004/UD=14022
         (c) 2004 American Chemical Society
*File 399: Use is subject to the terms of your user/customer agreement.
Alert feature enhanced for multiple files, etc. See HELP ALERT.
  File 434:SciSearch(R) Cited Ref Sci 1974-1989/Dec
         (c) 1998 Inst for Sci Info
  File
         8:Ei Compendex(R) 1970-2004/May W3
         (c) 2004 Elsevier Eng. Info. Inc.
        99: Wilson Appl. Sci & Tech Abs 1983-2004/Apr
         (c) 2004 The HW Wilson Co.
```

```
File 135:NewsRx Weekly Reports 1995-2004/May W3
         (c) 2004 NewsRx
*File 135: New newsletters are now added. See Help News135 for the
complete list of newsletters.
  File 266: FEDRIP 2004/Mar
         Comp & dist by NTIS, Intl Copyright All Rights Res
  File 315: ChemEng & Biotec Abs 1970-2004/Apr
         (c) 2004 DECHEMA
  File 357: Derwent Biotech Res. _1982-2004/May W4
         (c) 2004 Thomson Derwent & ISI
  File 358:Current BioTech Abs 1983-2004/Apr
         (c) 2004 DECHEMA
      Set Items Description
      ---
? s P2X(2)receptor
           0 P2X(2)RECEPTOR
? s P()2()X (n2) receptor
Processing
Processing
Processed 10 of 29 files ...
Processing
Processed 20 of 29 files ...
Processing
Completed processing all files
         8551901 P
        24271847 2
         5061576 X
         3785095 RECEPTOR
             10 P()2()X (N2) RECEPTOR
      S2
? s p2X (n2)receptor
           10179 P2X
         3785095 RECEPTOR
      S3
            5539 P2X (N2) RECEPTOR
? s "p2x" (n2) receptor
           10179 P2X
         3785095 RECEPTOR
            5539 "P2X" (N2) RECEPTOR
      S4
? s p2X4 (n2) receptor
             714 P2X4
         3785095 RECEPTOR
      S5
           277 P2X4 (N2) RECEPTOR
? s s3 (n2) antagonist
          5539 S3
841251 ANTAGONIST
            383 S3 (N2) ANTAGONIST
      S 6
? s s4 (n2)antagonist
          5539 S4
841251 ANTAGONIST
383 S4 (N2)ANTAGONIST
      S7
? ds
Set
        Items
                Description
S1
            0
                P2X(2) RECEPTOR
                P()2()X (N2) RECEPTOR
S2
           10
S3
         5539
                P2X (N2) RECEPTOR
                "P2X" (N2) RECEPTOR
S4
         5539
                P2X4 (N2) RECEPTOR
S_5
          277
S6
          383
                S3 (N2) ANTAGONIST
                S4 (N2)ANTAGONIST
S7
          383
```

? s s3 (n2) agonist

```
5539 S3
          542915 AGONIST
      S8
             390 S3 (N2) AGONIST
? s s4(n2)agonist
            5539
                 S4
          542915 AGONIST
      S9
             390 S4(N2)AGONIST
? ds
Set
        Items
                Description
S1
            0
                P2X(2) RECEPTOR
S2
           10
                P()2()X (N2) RECEPTOR
S3
         5539
                P2X (N2) RECEPTOR
                "P2X" (N2) RECEPTOR
         5539
S4
                P2X4 (N2) RECEPTOR
S5
          277
S6
          383
                S3 (N2) ANTAGONIST
Š7
          383
                S4 (N2) ANTAGONIST
S8
          390
                S3 (N2) AGONIST
          390
S9
                S4 (N2) AGONIST
? s s5 (n2) antagonist
             277 S5
          841251 ANTAGONIST
     S10
              2 S5 (N2) ANTAGONIST
? s s5 (N2)agonist
             277 S5
          542915 AGONIST
              15 S5 (N2)AGONIST
? s s10(n5)inhibit?
Processing
Processed 10 of 29 files ...
Completed processing all files
               2 S10
         6944746 INHIBIT?
              1 S10(N5)INHIBIT?
     S12
? s s11(n5)inhibit?
Processing
Processed 20 of 29 files ...
Completed processing all files
              15 S11
         6944746 INHIBIT?
               0 S11(N5)INHIBIT?
     S13
? t s12/3, k/
>>>KWIC option is not available in file(s): 399
             (Item 1 from file: 5)
 12/3, K/1
DIALOG(R) File
              5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
0013366531
             BIOSIS NO.: 200100538370
Mutation of histidine 241 of the rat P2X4 receptor alters agonist and
  antagonist sensitivities
AUTHOR: Xiong K (Reprint); Li C (Reprint); Stewart R R (Reprint); Weight F
  F (Reprint)
AUTHOR ADDRESS: Laboratory of Molecular and Cellular Neurobiology, NIAAA,
  NIH, Rockville, MD, USA**USA
JOURNAL: Society for Neuroscience Abstracts 27 (2): p1571 2001 2001
MEDIUM: print
CONFERENCE/MEETING: 31st Annual Meeting of the Society for Neuroscience
San Diego, California, USA November 10-15, 2001; 20011110
ISSN: 0190-5295
DOCUMENT TYPE: Meeting; Meeting Abstract
```

RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: 55 muM). In addition, the mutation of histidine 241 induced a sensitivity of the rat **P2X4 receptor** to the **antagonist**, PPADS: 100 muM PPADS **inhibited** 50 muM ATP-activated current by 5% in the wildtype and by 95% in the...

? ds

```
Set
        Items
                 Description
S1
            0
                 P2X(2) RECEPTOR
S2
           10
                 P()2()X (N2) RECEPTOR
S3
         5539
                 P2X (N2) RECEPTOR
                 "P2X" (N2) RECEPTOR
S4
         5539
          277
                P2X4 (N2) RECEPTOR
S5
          383
                S3 (N2) ANTAGONIST
S6
S7
          383
                 S4 (N2) ANTAGONIST
S8
          390
                 S3 (N2) AGONIST
S9
          390
                 S4 (N2) AGONIST
S10
            2
                S5 (N2) ANTAGONIST
                 S5 (N2) AGONIST
S11
           15
                 S10(N5) INHIBIT?
S12
            1
S13
            0
                 S11(N5)INHIBIT?
? t s10/3, k/all
>>>KWIC option is not available in file(s): 399
```

10/3,K/1 (Item 1 from file: 5)

DIALOG(R) File 5: Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0013366531 BIOSIS NO.: 200100538370

Mutation of histidine 241 of the rat P2X4 receptor alters agonist and antagonist sensitivities

AUTHOR: Xiong K (Reprint); Li C (Reprint); Stewart R R (Reprint); Weight F (Reprint)

AUTHOR ADDRESS: Laboratory of Molecular and Cellular Neurobiology, NIAAA, NIH, Rockville, MD, USA**USA

JOURNAL: Society for Neuroscience Abstracts 27 (2): p1571 2001 2001

MEDIUM: print

CONFERENCE/MEETING: 31st Annual Meeting of the Society for Neuroscience

San Diego, California, USA November 10-15, 2001; 20011110

ISSN: 0190-5295

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: 55 muM). In addition, the mutation of histidine 241 induced a sensitivity of the rat **P2X4 receptor** to the **antagonist**, PPADS: 100 muM PPADS inhibited 50 muM ATP-activated current by 5% in the wildtype...

10/3,K/2 (Item 1 from file: 399)

DIALOG(R) File 399:CA SEARCH(R)

(c) 2004 American Chemical Society. All rts. reserv.

139317806 CA: 139(21)317806j JOURNAL

Up-regulation of P2X2, P2X4 receptor and ischemic cell death: prevention by P2 antagonists

AUTHOR(S): Cavaliere, F.; Florenzano, F.; Amadio, S.; Fusco, F. R.; Viscomi, M. T.; D'Ambrosi, N.; Vacca, F.; Sancesario, G.; Bernardi, G.;

- look into.

Molinari, M.; Volonte, C.

LOCATION: IRCCS Santa Lucia Foundation, Rome, Italy

JOURNAL: Neuroscience (Oxford, U. K.) (Neuroscience (Oxford, United

Kingdom)) DATE: 2003 VOLUME: 120 NUMBER: 1 PAGES: 85-98 CODEN: NRSCDN

ISSN: 0306-4522 PUBLISHER ITEM IDENTIFIER: 0306-4522(03)00228-8

LANGUAGE: English PUBLISHER: Elsevier Science Ltd.

? t s11/3, k/all

>>>KWIC option is not available in file(s): 399

11/3,K/1 (Item 1 from file: 5)

DIALOG(R) File 5: Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0014336193 BIOSIS NO.: 200300294012

MUTATION OF HISTIDINES IN THE RAT P2X4 RECEPTOR ALTERS AGONIST POTENCY: GATING VERSUS BINDING.

AUTHOR: Xiong K (Reprint); Stewart R R (Reprint); Hu X Q (Reprint); Werby E (Reprint); Weight F F (Reprint); Li C (Reprint)

AUTHOR ADDRESS: Laboratory of Molecular and Cellular Neurobiology, NIAAA, NIH, Bethesda, MD, USA**USA

JOURNAL: Society for Neuroscience Abstract Viewer and Itinerary Planner 2002 pAbstract No. 337.4 2002 2002

MEDIUM: cd-rom

CONFERENCE/MEETING: 32nd Annual Meeting of the Society for Neuroscience

Orlando, Florida, USA November 02-07, 2002; 20021102

SPONSOR: Society for Neuroscience

DOCUMENT TYPE: Meeting; Meeting Poster; Meeting Abstract

RECORD TYPE: Abstract LANGUAGE: English

MUTATION OF HISTIDINES IN THE RAT P2X4 RECEPTOR ALTERS AGONIST POTENCY: GATING VERSUS BINDING.

...ABSTRACT: investigated the possibility that the three histidine residues in the extracellular loop of the rat **P2X4 receptor** regulate **agonist** potency. Mutation of histidine 241 to alanine (H241A) decreased the EC50 value of the ATP...

...competitive ATP binding curves were similar. These results suggest that histidine 241 of the rat **P2X4 receptor** regulates **agonist** potency by altering receptor gating rather than agonist binding.) K.X. and R.R.S...

11/3,K/2 (Item 2 from file: 5)

DIALOG(R) File 5:Biosis Previews(R) (c) 2004 BIOSIS. All rts. reserv.

0014141199 BIOSIS NO.: 200300099918

Multiple purinergic receptors lead to intracellular calcium increases in cultured rat Sertoli cells.

AUTHOR: Ko W H (Reprint); Au C L; Yip C Y

AUTHOR ADDRESS: Department of Physiology, Faculty of Medicine, Chinese University of Hong Kong, Shatin, Basic Medical Sciences Building, Hong Kong, China**China

AUTHOR E-MAIL ADDRESS: whko@cuhk.edu.hk

JOURNAL: Life Sciences 72 (13): p1519-1535 February 14, 2003 2003

MEDIUM: print

ISSN: 0024-3205 (ISSN print)

DOCUMENT TYPE: Article RECORD TYPE: Abstract

LANGUAGE: English

...ABSTRACT: one additional receptor population that allowed nucleotides to increase (Ca2+)i. Apart from the P2Y receptor agonists, the P2X4 and P2X7 agonist, 2' and 3'-O-(4-benzoylbenzoyl)-ATP (Bz-ATP), also evoked (Ca2+)i increases in...

11/3,K/3 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0013790313 BIOSIS NO.: 200200383824

P2X purinergic receptor channel expression and function in bovine aortic endothelium

AUTHOR: Ramirez Angelina N (Reprint); Kunze Diana L

AUTHOR ADDRESS: MetroHealth Systems, 2500 MetroHealth Drive, Rammelkamp

Center R334, Cleveland, OH, 44109-1998, USA**USA

JOURNAL: American Journal of Physiology 282 (6 Part 2): pH2106-H2116 June,

2002 2002

MEDIUM: print ISSN: 0002-9513

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: whole cell and outside-out patch recordings using 2-methyl-thio-ATP (MeSATP) as a **P2X4** and P2X5 **receptor agonist** and 2',3'-O-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3,K/4 (Item 4 from file: 5)

DIALOG(R) File 5: Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0013366531 BIOSIS NO.: 200100538370

Mutation of histidine 241 of the rat P2X4 receptor alters agonist and antagonist sensitivities

AUTHOR: Xiong K (Reprint); Li C (Reprint); Stewart R R (Reprint); Weight F (Reprint)

AUTHOR ADDRESS: Laboratory of Molecular and Cellular Neurobiology, NIAAA, NIH, Rockville, MD, USA**USA

JOURNAL: Society for Neuroscience Abstracts 27 (2): p1571 2001 2001

MEDIUM: print

CONFERENCE/MEETING: 31st Annual Meeting of the Society for Neuroscience

San Diego, California, USA November 10-15, 2001; 20011110

ISSN: 0190-5295

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract LANGUAGE: English

Mutation of histidine 241 of the rat P2X4 receptor alters agonist and antagonist sensitivities

11/3,K/5 (Item 5 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)

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0013092289 BIOSIS NO.: 200100264128

P2X receptors in bovine aortic endothelium

AUTHOR: Ramirez Angelina N (Reprint); Kunze Diana L (Reprint)

AUTHOR ADDRESS: Rammelkamp Center, MetroHealth, Case Western Reserve University, 2500 MetroHealth Drive, Cleveland, OH, 44109, USA**USA

JOURNAL: FASEB Journal 15 (4): pA109 March 7, 2001 2001

MEDIUM: print

CONFERENCE/MEETING: Annual Meeting of the Federation of American Societies for Experimental Biology on Experimental Biology 2001 Orlando, Florida, USA March 31-April 04, 2001; 20010331

ISSN: 0892-6638

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: electrophysiological effect of ligand activation of these P2X receptors we used ATP, 2Me-S-ATP (P2X4 and P2X5 receptor agonist) and Bz-ATP (P2X7 receptor agonist) to characterize the channels using whole cell and outside...

11/3,K/6 (Item 6 from file: 5)

DIALOG(R) File 5:Biosis Previews(R) (c) 2004 BIOSIS. All rts. reserv.

0012472736 BIOSIS NO.: 200000191049

Mutation of histidine 286 of the human P2X4 purinoceptor removes extracellular pH sensitivity

AUTHOR: Clarke C E; Benham C D (Reprint); Bridges A; George A R; Meadows H

AUTHOR ADDRESS: Department of Neuroscience, SmithKline Beecham

Pharmaceuticals, Third Avenue, Harlow, New Frontiers Science Park, Essex, CM19 5AW, UK**UK

JOURNAL: Journal of Physiology (Cambridge) 523 (3): p697-703 march 15, 2000 2000

MEDIUM: print

ISSN: 0022-3751

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: directed mutagenesis of histidine 286 (H286) to alanine completely abolished the pH sensitivity of the **P2X4 receptor** at all **agonist** concentrations. ATP potency showed a small (fourfold) leftward shift. Mutagenesis of the other three histidines...

11/3,K/7 (Item 1 from file: 34)

DIALOG(R) File 34: SciSearch(R) Cited Ref Sci (c) 2004 Inst for Sci Info. All rts. reserv.

10627774 Genuine Article#: 550YC No. References: 42

Title: P2X purinergic receptor channel expression and function in bovine aortic endothelium

Author(s): Ramirez AN (REPRINT) ; Kunze DL

Corporate Source: Metrohlth Syst,Rammelkamp Ctr R334,2500 Metrohlth Dr/Cleveland//OH/44109 (REPRINT); Metrohlth Syst,Rammelkamp Ctr Educ & Res,Cleveland//OH/44109; Case Western Reserve Univ,Dept

Neurosci, Cleveland//OH/44109 Journal: AMERICAN JOURNAL OF PHYSIOLOGY-HEART AND CIRCULATORY PHYSIOLOGY,

have

2002, V282, N6 (JUN), PH2106-H2116

ISSN: 0363-6135 Publication date: 20020600

Publisher: AMER PHYSIOLOGICAL SOC, 9650 ROCKVILLE PIKE, BETHESDA, MD 20814

USA

Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

...Abstract: whole cell and outside-out patch recordings using 2-methyl-thio-ATP (MeSATP) as a **P2X4** and P2X5 **receptor agonist** and 2',3'-O-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3,K/8 (Item 1 from file: 50)

DIALOG(R) File 50: CAB Abstracts

(c) 2004 CAB International. All rts. reserv.

04392110 CAB Accession Number: 20033016414

Multiple purinergic receptors lead to intracellular calcium increases in cultured rat Sertoli cells.

Ko, W. H.; Au, C. L.; Yip, C. Y.

Department of Physiology, Faculty of Medicine, The Chinese University of Hong Kong, Basic Medical Sciences Building, Shatin, Hong Kong, China.

Life Sciences vol. 72 (13): p.1519=1535

Publication Year: 2003 ISSN: 0024-3205 --Language: English

Document Type: Journal article

... one additional receptor population that allowed nucleotides to increase (Ca2+)i. Apart from the P2Y **receptor** agonists, the **P2X4** and P2X7 **agonist** , 2' and 3'-O-(4-benzoylbenzoyl)-ATP (Bz-ATP), also evoked (Ca2+)i increases in...

11/3,K/9 (Item 1 from file: 71)

DIALOG(R) File 71: ELSEVIER BIOBASE

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02061326 2002141757

P2X purinergic receptor channel expression and function in bovine aortic endothelium

Ramirez A.N.; Kunze D.L.

ADDRESS: A.N. Ramirez, Rammelkamp Center R334, MetroHealth Systems, 2500 MetroHealth Drive, Cleveland, OH 44109-1998, United States

EMAIL: arnavarro@metrohealth.org

Journal: American Journal of Physiology - Heart and Circulatory Physiology , 282/6 51-6 (H2106-H2116), 2002, United States

CODEN: AJPPD ISSN: 0363-6135

DOCUMENT TYPE: Article

LANGUAGES: English SUMMARY LANGUAGES: English

NO. OF REFERENCES: 42

...whole cell and outside-out patch recordings using 2-methyl-thio-ATP (MeSATP) as a **P2X4** and P2X5 **receptor agonist** and 2prime, 3prime-O-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3,K/10 (Item 1 from file: 73)

DIALOG(R) File 73: EMBASE

(c) 2004 Elsevier Science B.V. All rts. reserv.

11650245 EMBASE No: 2002220993

P2X purinergic receptor channel expression and function in bovine aortic endothelium

Ramirez A.N.; Kunze D.L.

A.N. Ramirez, Rammelkamp Center R334, MetroHealth Systems, 2500

MetroHealth Drive, Cleveland, OH 44109-1998 United States

AUTHOR EMAIL: arnavarro@metrohealth.org

American Journal of Physiology - Heart and Circulatory Physiology (AM. J. PHYSIOL. HEART CIRC. PHYSIOL.) (United States) 2002, 282/6 51-6

(H2106-H2116)

CODEN: AJPPD ISSN: 0363-6135 DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 42

...whole cell and outside-out patch recordings using 2-methyl-thio-ATP (MeSATP) as a **P2X4** and P2X5 **receptor agonist** and 2prime,3prime-O-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3,K/11 (Item 1 from file: 98)

DIALOG(R)File 98:General Sci Abs/Full-Text (c) 2004 The HW Wilson Co. All rts. reserv.

04873124 H.W. WILSON RECORD NUMBER: BGSA02123124

P2X purinergic receptor channel expression and function in bovine aortic endothelium.

Ramirez, Angelina

Kunze, Diana L

American Journal of Physiology (Am J Physiol) v. 282 no6 (June 2002 pt2) p. H2106-H2116

SPECIAL FEATURES: bibl graph il ISSN: 0002-9513

LANGUAGE: English

COUNTRY OF PUBLICATION: United States

...ABSTRACT: whole cell and outside-out patch recordings using 2-methyl-thio-ATP (MeSATP) as a **P2X4** and P2X5 lreceptor agonist and 2',3'-O-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3,K/12 (Item 1 from file: 144)

DIALOG(R) File 144: Pascal

(c) 2004 INIST/CNRS. All rts. reserv.

15708491 PASCAL No.: 02-0417633

P2X purinergic receptor channel expression and function in bovine aortic endothelium

RAMIREZ Angelina N; KUNZE Diana L

Rammelkamp Center for Education and Research, MetroHealth Systems and Department of Neurosciences, Case Western Reserve University, Cleveland, Ohio 44109-1998, United States

Journal: American journal of physiology. Heart and circulatory physiology, 2002, 51 (6) H2106-H2116

Language: English

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... whole cell and outside-out patch recordings using 2-methyl-thio-ATP (MeSATP) as P2X4 and P2X5 agonist а receptor 2',3'-O-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3,K/13 (Item 1 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

11814707 PMID: 12003818

P2X purinergic receptor channel expression and function in bovine aortic endothelium.

Ramirez Angelina N; Kunze Diana L

Rammelkamp Center for Education and Research, MetroHealth Systems and Department of Neurosciences, Case Western Reserve University, Cleveland, Ohio 44109-1998, USA. arnavarro@metrohealth.org

American journal of physiology. Heart and circulatory physiology (United Jun 2002, 282 (6) pH2106-16, ISSN 0363-6135 Journal Code: States) 100901228

Contract/Grant No.: HL-61436; HL; NHLBI

Document type: Journal Article

Languages: ENGLISH Main Citation Owner: NLM Record type: Completed

... whole cell and outside-out patch recordings using 2-methyl-thio-ATP P2X4 and P2X5 receptor (MeSATP) as a agonist 2',3'-0-(4-benzoylbenzoyl)ATP (BzATP) as a P2X7 receptor agonist. MeSATP (10...

11/3, K/14(Item 2 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

10612714 PMID: 10718748

Mutation of histidine 286 of the human P2X4 purinoceptor removes extracellular pH sensitivity.

Clarke C E; Benham C D; Bridges A; George A R; Meadows H J

Departments of Neuroscience, Biotechnology and Genetics and Medicinal Chemistry Research, SmithKline Beecham Pharmaceuticals, New Frontiers Science Park, Third Avenue, Harlow, Essex CM19 5AW, UK.

Journal of physiology (ENGLAND) Mar 15 2000, 523 Pt 3 p697-703, ISSN 0022-3751 Journal Code: 0266262

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

... directed mutagenesis of histidine 286 (H286) to alanine completely abolished the pH sensitivity of the P2X4 receptor at all agonist concentrations. ATP potency showed a small (fourfold) leftward shift. Mutagenesis of the other three histidines...

11/3,K/15 (Item 1 from file: 156)

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DIALOG(R) File 156: ToxFile
(c) format only 2004 The Dialog Corporation. All rts. reserv.
            NLM Doc No: 10718748
  Mutation of histidine 286 of the human P2X4 purinoceptor removes
extracellular pH sensitivity.
  Clarke C E; Benham C D; Bridges A; George A R; Meadows H J
Departments of Neuroscience, Biotechnology and Genetics and Medicinal Chemistry Research, SmithKline Beecham Pharmaceuticals, New Frontiers
Science Park, Third Avenue, Harlow, Essex CM19 5AW, UK. Journal Name: Journal of physiology (ENGLAND) Pub.
                                                       Pub. Year: Mar 15 2000
523 Pt 3 p697-703, ISSN: 0022-3751 Journal Code: 0266262
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: Completed
  ... directed mutagenesis of histidine 286 (H286) to alanine completely
abolished the pH sensitivity of the P2X4
                                                      receptor at all agonist
(fourfold) leftward shift.
concentrations. ATP potency showed a small Mutagenesis of the other three histidines...
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S1
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S2
            10
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S3
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                  P2X (N2) RECEPTOR
                 "P2X" (N2) RECEPTOR
S4
          5539
S5
           277
                  P2X4 (N2) RECEPTOR
                  S3 (N2) ANTAGONIST
S 6
           383
S7
           383
                  S4 (N2) ANTAGONIST
S8
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                  S3 (N2) AGONIST
S9
           390
                  S4 (N2) AGONIST
S10
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                  S5 (N2) ANTAGONIST
                  S5 (N2) AGONIST
S11
            15
S12
             1
                  S10 (N5) INHIBIT?
S13
             0
                  S11(N5)INHIBIT?
? neuropathic (n2)pain
>>>Unrecognizable Command
? s neuropathic(n2)pain
            32882 NEUROPATHIC
          1035947 PAIN
            19158 NEUROPATHIC (N2) PAIN
     S14
? s tactile(n2)allodynia
            46026 TACTILE
            11269
                   ALLODYNIA
             1570 TACTILE (N2) ALLODYNIA
     S15
? s s14 and s15
            19158
                    S14
             1570
                   S15
              932 S14 AND S15
     S16
...examined 50 records
...examined 50 records
                          (100)
...examined 50 records
                          (150)
...examined 50 records
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...examined 50 records
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...examined 50 records (450)
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...examined 50 records
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...examined 50 records
                        (900)
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                Description
Set
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S2
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                P2X (N2) RECEPTOR
S3
                "P2X" (N2) RECEPTOR
S4
         5539
          277
                P2X4 (N2) RECEPTOR
S5
          383
                S3 (N2) ANTAGONIST
S 6
                S4 (N2) ANTAGONIST
S7
          383
S8
          390
                S3 (N2) AGONIST
          390
                S4 (N2) AGONIST
S 9
           2
                S5 (N2) ANTAGONIST
S10
S11
           15
                S5 (N2) AGONIST
S12
            1
                S10(N5)INHIBIT?
S13
            0
                S11(N5)INHIBIT?
                NEUROPATHIC (N2) PAIN
S14
        19158
         1570
                TACTILE (N2) ALLODYNIA
S15
                S14 AND S15
          932
S16
          344
                RD (unique items)
S17
? s s17 and s5
             344 S17
             277
                  S5
               2 S17 AND S5
     S18
? t/3, k/all
>>>KWIC option is not available in file(s): 399
 18/3,K/1
              (Item 1 from file: 5)
                5:Biosis Previews(R)
DIALOG(R) File
(c) 2004 BIOSIS. All rts. reserv.
            BIOSIS NO.: 200400233892
0014864183
Up-regulation of microglial P2X4 receptor expression by retinoic acid.
AUTHOR: Tozaki Hidetoshi (Reprint); Koizumi Shuichi; Inoue Kazuhide
  (Reprint)
AUTHOR ADDRESS: Div. Biosignal, Div. Pharmacol., Natl. Inst. Hlth. Sci.,
  Tokyo, 158-8501, Japan**Japan
JOURNAL: Journal of Pharmacological Sciences 94 (Supplement 1): p299P 2004
 2004
MEDIUM: print
CONFERENCE/MEETING: 77th Annual Meeting of the Japanese Pharmacological
Society Osaka, Japan March 08-10, 2004; 20040308
SPONSOR: Japanese Pharmacological Society
ISSN: 1347-8613 (ISSN print)
DOCUMENT TYPE: Meeting; Meeting Abstract
RECORD TYPE: Citation
```

LANGUAGE: English

Up-regulation of microglial P2X4 receptor expression by retinoic acid.

DESCRIPTORS:

...DISEASES: neuropathic pain --...

... tactile allodynia --

18/3,K/2 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.

0014790941 BIOSIS NO.: 200400171698

ATP- and adenosine-mediated signaling in the central nervous system:
Chronic pain and microglia: Involvement of the ATP receptor P2X4.

AUTHOR: Inoue Kazuhide (Reprint); Tsuda Makoto; Koizumi Schuichi
AUTHOR ADDRESS: Division of Biosignaling, National Institute of Health
Sciences, 1-18-1 Kamiyoga, Setagaya-ku, Tokyo, 158-8501, Japan**Japan
AUTHOR E-MAIL ADDRESS: inoue@nihs.go.jp

MEDIUM: print

ISSN: 1347-8613 _(ISSN print)

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

- ...mediated signaling in the central nervous system: Chronic pain and microglia: Involvement of the ATP receptor P2X4 .
- ...ABSTRACT: reported that activation of P2X2/3 heteromeric channel/receptor in primary sensory neurons causes acutely tactile allodynia, one hallmark of neuropathic pain. We report here that tactile allodynia under the chronic pain state requires an activation of the P2X4 ionotropic ATP receptor and p38 mitogen-activated protein kinase (MAPK) in spinal cord microglia. Two weeks after L5...
- ...results demonstrate that activation of P2X4 or p38MAPK in spinal cord microglia is necessary for **tactile allodynia** after nerve injury. DESCRIPTORS:

MISCELLANEOUS TERMS: tactile allodynia ? ds

Set	Items	Description
S1	0	P2X (2) RECEPTOR
S2	10	P()2()X (N2) RECEPTOR
s3	5539	P2X (N2) RECEPTOR
S4	5539	"P2X" (N2) RECEPTOR
S5	277	P2X4 (N2) RECEPTOR
S6	383	S3 (N2) ANTAGONIST
s7	383	S4 (N2)ANTAGONIST
S8	390	S3 (N2) AGONIST
S9	390	S4 (N2) AGONIST
S10	2	S5 (N2) ANTAGONIST
S11	15	S5 (N2)AGONIST
S12	1	S10 (N5) INHIBIT?
S13	0	S11 (N5) INHIBIT?
S14	19158	NEUROPATHIC (N2) PAIN
S15	1570	TACTILE (N2) ALLODYNIA
S16	932	S14 AND S15

```
S17
          344
                RD (unique items)
            2
                S17 AND S5
S18
? s 17 and s4
         2491709
                 17
            5539
                 S4
             148 17 AND S4
     S19
...examined 50 records (50)
...examined 50 records (100)
...completed examining records
     S20
              76 RD (unique items)
? s s17 and s4
             344
                 S17
            5539 S4
               5 S17 AND S4
     S21
? rd
...completed examining records
     S22
               5 RD (unique items)
? t/3, k/all
>>>KWIC option is not available in file(s): 399
 22/3,K/1
              (Item 1 from file: 5)
DIALOG(R) File
                5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
             BIOSIS NO.: 200400233892
Up-regulation of microglial P2X4 receptor expression by retinoic acid.
AUTHOR: Tozaki Hidetoshi (Reprint); Koizumi Shuichi; Inoue Kazuhide
  (Reprint)
AUTHOR ADDRESS: Div. Biosignal, Div. Pharmacol., Natl. Inst. Hlth. Sci.,
  Tokyo, 158-8501, Japan**Japan
JOURNAL: Journal of Pharmacological Sciences 94 (Supplement 1): p299P 2004
 2004
MEDIUM: print
CONFERENCE/MEETING: 77th Annual Meeting of the Japanese Pharmacological
Society Osaka, Japan March 08-10, 2004; 20040308
SPONSOR: Japanese Pharmacological Society
ISSN: 1347-8613 _(ISSN print)
DOCUMENT TYPE: Meeting; Meeting Abstract
RECORD TYPE: Citation
LANGUAGE: English
DESCRIPTORS:
  ...DISEASES: neuropathic
                              pain --...
... tactile
              allodynia --
  CHEMICALS & BIOCHEMICALS:
                              ... P2X -4 receptor --
 22/3,K/2
              (Item 2 from file: 5)
DIALOG(R) File
               5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
0014790941
             BIOSIS NO.: 200400171698
ATP- and adenosine-mediated signaling in the central nervous system:
  Chronic pain and microglia: Involvement of the ATP receptor P2X4.
AUTHOR: Inoue Kazuhide (Reprint); Tsuda Makoto; Koizumi Schuichi
AUTHOR ADDRESS: Division of Biosignaling, National Institute of Health
  Sciences, 1-18-1 Kamiyoga, Setagaya-ku, Tokyo, 158-8501, Japan**Japan
AUTHOR E-MAIL ADDRESS: inoue@nihs.go.jp
JOURNAL: Journal of Pharmacological Sciences 94 (2): p112-114 February
```

2004 2004 MEDIUM: print

ISSN: 1347-8613 _(ISSN print)

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: reported that activation of P2X2/3 heteromeric channel/receptor in primary sensory neurons causes acutely tactile allodynia, one hallmark of neuropathic pain. We report here that tactile allodynia under the chronic pain state requires an activation of the P2X4 ionotropic ATP receptor and...

...results demonstrate that activation of P2X4 or p38MAPK in spinal cord microglia is necessary for **tactile allodynia** after nerve injury. DESCRIPTORS:

CHEMICALS & BIOCHEMICALS: ... P2X -4 receptor -- MISCELLANEOUS TERMS: tactile allodynia

22/3,K/3 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.

0014723989 BIOSIS NO.: 200400092758

Effects of A-317491, a novel and selective P2X3/P2X2/3 receptor antagonist, on neuropathic, inflammatory and chemogenic nociception following intrathecal and intraplantar administration.

AUTHOR: McGaraughty Steve (Reprint); Wismer Carol T; Zhu Chang Z; Mikusa Joseph; Honore Prisca; Chu Katharine L; Lee Chih-Hung; Faltynek Connie R; Jarvis Michael F

AUTHOR ADDRESS: Neuroscience Research, Global Pharmaceutical Research and Development, Abbott Laboratories, 100 Abbott Park Road, R4PM, AP9-1, Abbott Park, IL, 60064-6118, USA**USA

AUTHOR E-MAIL ADDRESS: Steve.P.Mcgaraughty@abbott.com

JOURNAL: British Journal of Pharmacology 140 (8): p1381-1388 December 2003 2003

MEDIUM: print

ISSN: 0007-1188 (ISSN print)

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: homomeric and P2X2/3 heteromeric channels, is antinociceptive in rat models of chronic inflammatory and **neuropathic pain**. In an effort to further evaluate the role of P2X3/P2X2/3 receptors in nociceptive...

...formalin assay (intrathecal ED50 = 10 nmol, intraplantar ED50 > 300 nmol). Nocifensive behaviors induced by the **P2X receptor** agonist alpha, beta-meATP were also significantly reduced by intraplantar injection of A-317491. 5...

...Intrathecal administration of A-317491 appears to be more effective than intraplantar administration to reduce tactile allodynia following peripheral nerve injury.

DESCRIPTORS:

...DISEASES: neuropathic pain --CHEMICALS & BIOCHEMICALS: ... P2X -2/3 receptor; ...

... P2X -3 receptor

60/4 in

22/3,K/4 (Item 1 from file: 34)
DIALOG(R)File 34:SciSearch(R) Cited Ref Sci

(c) 2004 Inst for Sci Info. All rts. reserv.

11245850 Genuine Article#: 627GF No. References: 35

Title: Alteration of dorsal root ganglion P2X (3) receptor expression and function following spinal nerve ligation in the rat

Corporate Source: Abbott Labs, Global Pharmaceutical Res & Dev, 100 Abbott Pk Rd/Abbott Pk//IL/60064 (REPRINT); Abbott Labs, Global Pharmaceutical Res & Dev, Abbott Pk//IL/60064

Journal: EXPERIMENTAL BRAIN RESEARCH, 2002, V147, N4 (DEC), P511-519

ISSN: 0014-4819 Publication date: 20021200

Publisher: SPRINGER-VERLAG, 175 FIFTH AVE, NEW YORK, NY 10010 USA Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

Title: Alteration of dorsal root ganglion P2X (3) receptor expression and function following spinal nerve ligation in the rat

Abstract: One subtype of ATP-gated ion channel, the P2X (3) receptor, is expressed primarily on peripheral sensory neurons. While it is known that P2X(3) receptors can participate in certain forms of nociceptive signaling, their involvement in neuropathic pain transmission is not known. We have examined the expression and function of P2X3 receptors in a rat spinal nerve ligation model of neuropathic pain. Fourteen days following L5/L6 spinal nerve ligation, the corresponding dorsal root ganglia (DRG) were...

...these were studied using immunohistochemical and electrophysiological techniques. Using a polyclonal antibody to label the P2X (3) receptor , a significant reduction in neuronal P2X(3) immunoreactivity was observed in the ipsilateral (injured) L5...

...Identifiers--SENSORY NEURONS; PURINERGIC SENSITIVITY; PERIPHERAL NEUROPATHY; **TACTILE ALLODYNIA**; ATP; INJURY; PAIN; NOCICEPTION; CURRENTS; BEHAVIOR

22/3,K/5 (Item 1 from file: 71)

DIALOG(R) File 71: ELSEVIER BIOBASE

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02545976 2004014674

Effects of A-317491, a novel and selective P2XSUB3/ P2X SUB2/3 receptor antagonist, on neuropathic, inflammatory and chemogenic nociception following intrathecal and intraplantar administration

McGaraughty S.; Wismer C.T.; Zhu C.Z.; Mikusa J.; Honore P.; Chu K.L.; Lee C.-H.; Faltynek C.R.; Jarvis M.F.

ADDRESS: S. McGaraughty, Neuroscience Research, Global Pharmaceutical R. and D., Abbott Laboratories, 100 Abbott Park Road, Abbott Park, IL 60064-6118, United States

EMAIL: Steve.P.Mcgaraughty@abbott.com

Journal: British Journal of Pharmacology, 140/8 (1381-1388), 2003, United Kingdom

CODEN: BJPCB

ISSN: 0007-1188

DOCUMENT TYPE: Article

LANGUAGES: English SUMMARY LANGUAGES: English

NO. OF REFERENCES: 48

Effects of A-317491, a novel and selective P2XSUB3/ P2X SUB2/3 receptor antagonist, on neuropathic, inflammatory and chemogenic nociception following intrathecal and intraplantar administration

...homomeric and P2XSUB2/3 heteromeric channels, is antinociceptive in rat models of chronic inflammatory and **neuropathic pain**. In an effort to further evaluate the role of P2XSUB3/P2XSUB2/3 receptors in nociceptive...

...formalin assay (intrathecal EDSUB50 = 10 nmol, intraplantar EDSUB50 > 300 nmol). Nocifensive behaviors induced by the **P2X receptor** agonist alpha, beta-meATP were also significantly reduced by intraplantar injection of A-317491. 5...

...Intrathecal administration of A-317491 appears to be more effective than intraplantar administration to reduce **tactile allodynia** following peripheral nerve injury.
? ds

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Set
        Items
                 Description
S1
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                 P()2()X (N2) RECEPTOR
S3
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S4
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                 "P2X" (N2) RECEPTOR
S5
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                 P2X4 (N2) RECEPTOR
S6
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                 S3 (N2) ANTAGONIST
S7
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S8
          390
                 S3 (N2) AGONIST
S9
          390
                S4 (N2) AGONIST
S10
                S5 (N2) ANTAGONIST
            2
S11
           15
                S5 (N2) AGONIST
S12
                S10 (N5) INHIBIT?
            1
S13
            0
                S11 (N5) INHIBIT?
        19158
S14
                NEUROPATHIC (N2) PAIN
S15
         1570
                TACTILE (N2) ALLODYNIA
S16
          932
                S14 AND S15
S17
          344
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                S17 AND S5
S18
            2
S19
                17 AND S4
          148
S20
           76
                RD (unique items)
S21
            5
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S22
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          826448 FLUX
           10506 ION(W)FLUX
     S23
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? s s4 and ion()flux
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                  ION
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          826448
                  FLUX
           10506
                  ION(W)FLUX
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? rd
...completed examining records
               5 RD (unique items)
? t/3, k/all
>>>KWIC option is not available in file(s): 399
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25/3,K/1 (Item 1 from file: 5)

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DIALOG(R) File
                5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
             BIOSIS NO.: 200200139889
0013546378
Potent P2X7 receptor antagonists: Tyrosyl derivatives synthesized using a
  sequential parallel synthetic approach
AUTHOR: Ravi R Gnana; Kertesy Sylvia B; Dubyak George R; Jacobson Kenneth A
  (Reprint)
AUTHOR ADDRESS: Molecular Recognition Section, LBC, NIH, NIDDK, Bldg. 8A,
  Rm. B1A-19, Bethesda, MD, 20892-0810, USA**USA
-JOURNAL: Drug-Development_Research_54-(2): p75-87-October, 2001-2001
MEDIUM: print
ISSN: 0272-4391
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
... ABSTRACT: MRS2306), displayed an IC50 value of 40 nM as an antagonist of
  P2X7 receptor-mediated ion
                               flux and was more potent than the
  reference compound 1. Nalpha-Cbz, Boc-piperazinyl derivatives, 11...
DESCRIPTORS:
  CHEMICALS & BIOCHEMICALS: ... P2X -7 receptor antagonists
 25/3,K/2
              (Item 2 from file: 5)
DIALOG(R)File
                5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
             BIOSIS NO.: 200100350446
0013178607
Structure-activity relationships of pyridoxal phosphate derivatives as
  potent and selective antagonists of P2X1 receptors
AUTHOR: Kim Yong-Chul; Brown Sean G; Harden T Kendall; Boyer Jose L; Dubyak
  George; King Brian F; Burnstock Geoffrey; Jacobson Kenneth A (Reprint)
AUTHOR ADDRESS: NIDDK, LBC, NIH, Bldg. 8A, Rm. B1A-19, Bethesda, MD,
  20892-0810, USA**USA
JOURNAL: Journal of Medicinal Chemistry 44 (3): p340-349 Debouary 1, 2001
2001
MEDIUM: print
ISSN: 0022-2623
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
... ABSTRACT: in functional assays at recombinant rat P2X1, P2X2, and P2X3
  receptors expressed in Xenopus oocytes ( ion
                                                 flux stimulation) and at
  turkey erythrocyte P2Y1 receptors (phospholipase C activation). Selected
  compounds were also evaluated as antagonists of ion
                                                        flux and the
  opening of a large pore at the recombinant human P2X7 receptor.
  Modifications were...
DESCRIPTORS:
  CHEMICALS & BIOCHEMICALS:
                               P2X -1 receptor; ...
... P2X -2 receptor ; ...
... P2X -3 receptor ; ...
... P2X -7 receptor; ...
... P2X -1 receptor antagonist agent, structure-activity relationships,
    synthesis
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25/3,K/3
              (Item 1 from file: 73)
DIALOG(R) File 73: EMBASE
(c) 2004 Elsevier Science B.V. All rts. reserv.
            EMBASE No: 2002018334
  Potent P2XSUB7 receptor antagonists: Tyrosyl derivatives synthesized
using a sequential parallel synthetic approach
  Ravi R.G.; Kertesy S.B.; Dubyak G.R.; Jacobson K.A.
  K.A. Jacobson, Molecular Recognition Section, NIH, NIDDK, Bethesda, MD
  20892-0810 United States
 AUTHOR EMAIL: kajacobs@helix.nih.gov
  Drug Development Research ( DRUG DEV. RES. ) (United States)
                                                                 2001, 54/2
  (75 - 87)
  CODEN: DDRED
                ISSN: 0272-4391
  DOCUMENT TYPE: Journal ; Article
  LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
 NUMBER OF REFERENCES: 32
  ...MRS2306), displayed an IC50 value of 40 nM as an antagonist of P2XSUB7
receptor-mediated ion flux and was more potent than the reference
compound 1. NSUPalpha-Cbz, Boc-piperazinyl derivatives, 11...
DRUG DESCRIPTORS:
              receptor; *purinergic receptor blocking agent--drug
*purine P2X
analysis--an; *purinergic receptor blocking agent--drug comparison--cm; *
purinergic receptor blocking...
              (Item 2 from file: 73)
 25/3,K/4
DIALOG(R) File 73: EMBASE
(c) 2004 Elsevier Science B.V. All rts. reserv.
             EMBASE No: 2001048256
11057087
  Structure-activity relationships of pyridoxal phosphate derivatives as
potent and selective antagonists of P2XSUB1 receptors
  Kim Y .- C.; Brown S.G.; Harden T.K.; Boyer J.L.; Dubyak G.; King B.F.;
Burnstock G.; Jacobson K.A.
  Dr. K.A. Jacobson, MRS, NIH, NIDDK, Bethesda, MD 20892-0810 United
  States
  AUTHOR EMAIL: kajacobs@helix.nih.gov
  Journal of Medicinal Chemistry ( J. MED. CHEM. ) (United States)
                                                                   01 FEB
  2001, 44/3 (340-349)
               ISSN: 0022-2623
  CODEN: JMCMA
  DOCUMENT TYPE: Journal ; Article
  LANGUAGE: ENGLISH
                     SUMMARY LANGUAGE: ENGLISH
  NUMBER OF REFERENCES: 43
  ...in functional assays at recombinant rat P2XSUB1, P2XSUB2, and P2XSUB3
receptors expressed in Xenopus oocytes ( ion flux stimulation) and at
turkey erythrocyte P2YSUB1 receptors (phospholipase C activation). Selected
compounds were also evaluated as antagonists of ion
                                                      flux and the
opening of a large pore at the recombinant human P2XSUB7 receptor.
Modifications were...
DRUG DESCRIPTORS:
...*drug analysis--an; *pyridoxal 5 phosphate--drug development--dv; *
pyridoxal 5 phosphate--pharmacology--pd; *purine P2X
                                                        receptor
--endogenous compound--ec
```

(Item 1 from file: 370) 25/3,K/5 DIALOG(R) File 370: Science (c) 1999 AAAS. All rts. reserv. (USE 9 FOR FULLTEXT) 00500325 The Cytolytic P.inf(2Z) Receptor for Extracellular ATP Identified as a P.inf(2X) Receptor (P2X .inf(7))

Surprenant, A.; Rassendren, F.; Kawashima, E.; North, R. A.; Buell, G. Glaxo Institute for Molecular Biology, Plan-les-Ouates, 1228 Geneva, Switzerland.

Science_Vol._272_5262_pp._7353

Publication Date: 5-03-1996 (960503) Publication Year: 1996

Document Type: Journal ISSN: 0036-8075

Language: English

Section Heading: Reports

Word Count: 2485

(THIS IS THE FULLTEXT)

The Cytolytic P.inf(2Z) Receptor for Extracellular ATP Identified as a P.inf(2X) Receptor (P2X .inf(7)) ...

...Abstract: channels, the P.inf(2X) receptors, are permeable only to small cations. Here, an ATP receptor , the P2X .inf(7) receptor , was cloned from rat brain and exhibited both these properties. This protein is homologous to...

Here, we isolated a P2X receptor complementary DNA (cDNA) ...Text: (P2X.inf(7)) that encodes a 595-amino acid protein (Fig. 1...

- ...two transmembrane domains, and a large extracellular loop (B8) . The COOH-terminal domain of the P2X .inf(7) receptor was much longer than that found in the other receptors but contained no further hydrophobic...
- ...span the membrane and showed no sequence homology with known proteins. The mRNA for the P2X .inf(7) receptor was strongly expressed in J774 and P815 macrophages, in microglia, brain, spinal cord, lung, and...
- ...to 2 s) of ATP evoked inward currents in HEK 293 cells into which the P2X .inf(7) receptor was transiently or stably transfected (Fig. 1B) (B10) . The agonist order of potency was BzATP...
- ... C and D). Antagonists had similar effects on J774 cells and HEK cells expressing the P2X .inf(7) receptor : currents evoked by 30 (mu) M BzATP were relatively insensitive to the purinoceptor antagonist suramin...
- ...inf(2Z) receptors (B2) , was ineffective (100 (mu) M, n = 6) at blocking current. The $\,$ P2X .inf(7) $\,$ receptor $\,$ thus presents a pharmacological profile typical of the receptor previously termed P.inf(2Z) (B1The P.inf(2Z) receptor has been characterized primarily by ion flux and dve uptake studies in macrophage-derived cell lines such as J774, particularly with the...
- ...4C) and did not cause uptake of YO-PRO-1 by HEK cells expressing the P2X .inf(2) receptor (Fig. 4, D through F). We tested the hypothesis that the unique COOH-terminal domain conferred these properties by repeating the experiments on HEK cells expressing the P2X .inf(7) receptor truncated to 418 amino acids (P2X.inf(7) (Delta) C, Fig. 1A). Agonist and antagonist
- ... Thus, the expression of a single protein, the P2X .inf(7) receptor,

endows cells with two distinct responses to ATP and its analog BzATP. The first, a...

- ...with other proteins intrinsic to HEK 293 cells might confer the cytolytic properties of the **P2X** .inf(7) **receptor** (B2) . Any such protein must be ubiquitous because we obtained qualitatively similar results from Chinese...
- ...receptor as a member of the P.inf(2X) family. The dual function of the P2X .inf(7) receptor, whereby it can operate both as an ion channel selective for small cations and inFigure F1 Caption: (A) Predicted amino acid sequence of the P2X .inf(7) receptor, aligned with that of the P2X .inf(2) receptor (B22). The middle line shows common amino acids, lines over the top sequence indicate probable...
- ...ATP and ATP analogs (as indicated) in low divalent solution obtained from cells expressing the P2X .inf(7) receptor . (C and D) Similar experiments on J774 cells; low divalent solutions also increased both amplitude...
- ...BzATP induced a sustained nonselective conductance. In (D), currents were recorded from HEK cells expressing **P2X** .inf(7) **receptor** in response to four 1-s applications of BzATP, with an interval of 12 min...2), or no receptors (Untrans.). BzATP concentration was 30 (mu) M for experiments with the **P2X** .inf(7) **receptor** and 300 (mu) M for all others (n = 6 throughout...

References and Notes:

...RACE-PCR; Life Technologies, Bethesda, MD) using poly(A).sup(+) RNA from the medial habenula. P2X .inf(7) receptor -specific sequences were amplified with two rounds of nested PCR, for which sense primers wereCCACGCGTCGACTAGTACGGGIIGGGIIGGGIIGand...

? ds

Set S1 S2		Items 0 10	Description P2X(2)RECEPTOR P()2()X (N2) RECEPTOR
S3		5539	** ** **
			P2X (N2) RECEPTOR
S4		5539	"P2X" (N2) RECEPTOR
S5		277	P2X4 (N2) RECEPTOR
S6		383	S3 (N2) ANTAGONIST
s7		383	S4 (N2) ANTAGONIST
S8		390	S3 (N2) AGONIST
S9		390	S4 (N2) AGONIST
S10		2	S5 (N2) ANTAGONIST
S11		15	S5 (N2)AGONIST
S12		1	S10(N5)INHIBIT?
S13		0	S11 (N5) INHIBIT?
S14		19158	NEUROPATHIC (N2) PAIN
S15		1570	TACTILE (N2) ALLODYNIA
S16		932	S14 AND S15
S17		344	RD (unique items)
S18		2	S17 AND S5
S19		148	17 AND S4
S20		76	RD (unique items)
S21		5	S17 AND S4
S22		5	RD (unique items)
S23		Ő	S5 AND ION()FLUX
S24			S4 AND ION()FLUX
S25		5	RD (unique items)
323 ? s	~ O	and inhib	
: 5	59	and Infill	OTC:

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Processing
 Processed 10 of 29 files ...
 Completed processing all files
              390 S9
          6944746 INHIBIT?
      S26
              194 S9 AND INHIBIT?
 ? s s9 (N5) inhibit?
 Processing
 Processed 20 of 29 files ...
 Completed processing all files
              390 S9
          6944746 INHIBIT?
                5 S9 (N5) INHIBIT?
      S27
 ? t/3, k/all
 >>>KWIC option is not available in file(s): 399
               (Item 1 from file: 5)
  27/3,K/1
 DIALOG(R) File
                5:Biosis Previews(R)
 (c) 2004 BIOSIS. All rts. reserv.
            BIOSIS NO.: 200000447266
 0012728953
 Influence of purinoceptor antagonism on diadenosine pentaphosphate-induced
   hypotension in anesthetized rats
 AUTHOR: Steinmetz Martin; Van Le Truc; Hollah Peter; Gabrieels Gert; Hohage
   Helge; Rahn Karl Heinz; Schlatter Eberhard (Reprint)
 AUTHOR ADDRESS: Medizinische Poliklinik, Experimentelle Nephrologie,
   Westfaelische Wilhelms-Universitaet, Domagkstrasse 3a, 48149, Muenster,
Germany**Germany

**JOURNAL: Journal of Pharmacology and Experimental Therapeutics 294 (3): p
 963-968 September, 2000 2000
 MEDIUM: print
 ISSN: 0022-3565
 DOCUMENT TYPE: Article
 RECORD TYPE: Abstract
 LANGUAGE: English
 ... ABSTRACT: the A2 purinoceptor antagonist
   3,7-dimethyl-1-propargylxanthine. The hypertensive effect by the
   prototype P2X
                   receptor
                               agonist alphabeta-methylene ATP was
   inhibited by pyridoxal-phosphate-6-azophenyl-2',4'-disulfonic acid, too.
   Purinoceptor antagonists reduced the maximal...
  27/3, K/2
               (Item 1 from file: 34)
 DIALOG(R) File 34:SciSearch(R) Cited Ref Sci
 (c) 2004 Inst for Sci Info. All rts. reserv.
 08949362
            Genuine Article#: 347QH
                                      No. References: 25
 Title: Influence of purinoceptor antagonism on diadenosine
     pentaphosphate-induced hypotension in anesthetized rats
 Author(s): Steinmetz M; Le VT; Hollah P; Gabriels G; Hohage H; Rahn KH;
     Schlatter E (REPRINT)
 Corporate Source: UNIV MUNSTER, MED POLIKLIN, DOMAGKSTR 3A/D-48149
     MUNSTER//GERMANY/ (REPRINT); UNIV MUNSTER, MED POLIKLIN/D-48149
     MUNSTER//GERMANY/
 Journal: JOURNAL OF PHARMACOLOGY AND EXPERIMENTAL THERAPEUTICS, 2000, V294
  N3 (SEP), P963-968
 ISSN: 0022-3565
                   Publication date: 20000900
 Publisher: AMER SOC PHARMACOLOGY EXPERIMENTAL THERAPEUTICS, 9650 ROCKVILLE
     PIKE, BETHESDA, MD 20814-3998
                                               (ABSTRACT AVAILABLE)
 Language: English Document Type: ARTICLE
```

... Abstract: A(2) purinoceptor antagonist 3,7-dimethyl-1-propargylxanthine, The hypertensive effect by the prototype P2X receptor alpha beta-methylene ATP was inhibited by pyridoxal-phosphate-6-azophenyl-2',4'-disulfonic acid, too. Purinoceptor antagonists reduced the maximal... 27/3,K/3 (Item 1 from file: 71) DIALOG(R) File 71: ELSEVIER BIOBASE (c) 2004 Elsevier Science B.V. All rts. reserv. 01521425 2000200567 Influence of purinoceptor antagonism on diadenosine pentaphosphate-induced hypotension in anesthetized rats Steinmetz M.; Le T.V.; Hollah P.; Gabriels G.; Hohage H.; Rahn K.H.; Schlatter E. ADDRESS: Dr. E. Schlatter, Medizinische Poliklinik, Experimentelle Nephrologie, Westfalische Wilhelms-Universitat, Domagkstrasse 3a, 48149 Munster, Germany EMAIL: eberhard.schlatter@uni-muenster.de Journal: Journal of Pharmacology and Experimental Therapeutics, 294/3 (963-968), 2000, United States CODEN: JPETA ISSN: 0022-3565 DOCUMENT TYPE: Article LANGUAGES: English SUMMARY LANGUAGES: English NO. OF REFERENCES: 25 ...Ainf 2 purinoceptor antagonist 3,7-dimethyl-1-propargylxanthine. The hypertensive effect by the prototype P2X receptor agonist alphabeta-methylene ATP was inhibited by pyridoxal-phosphate-6-azophenyl-2',4'-disulfonic acid, too. Purinoceptor antagonists reduced the maximal... 27/3,K/4 (Item 1 from file: 73) DIALOG(R) File 73: EMBASE (c) 2004 Elsevier Science B.V. All rts. reserv. EMBASE No: 2000309756 Influence of purinoceptor antagonism on diadenosine pentaphosphate-induced hypotension in anesthetized rats Steinmetz M.; Le T.V.; Hollah P.; Gabriels G.; Hohage H.; Rahn K.H.; Schlatter E. Dr. E. Schlatter, Medizinische Poliklinik, Experimentelle Nephrologie, Westfalische Wilhelms-Universitat, Domagkstrasse 3a, 48149 Munster Germany AUTHOR EMAIL: eberhard.schlatter@uni-muenster.de Journal of Pharmacology and Experimental Therapeutics (J. PHARMACOL. EXP. THER.) (United States) 2000, 294/3 (963-968) CODEN: JPETA ISSN: 0022-3565 DOCUMENT TYPE: Journal; Article LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH NUMBER OF REFERENCES: 25 ... Ainf 2 purinoceptor antagonist 3,7-dimethyl-1-propargylxanthine. The hypertensive effect by the prototype P2X receptor

pyridoxal-phosphate-6-azophenyl-2',4'-disulfonic acid, too. Purinoceptor

alphabeta-methylene ATP was inhibited by

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antagonists reduced the maximal...
 27/3,K/5
              (Item 1 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
10820877
          PMID: 10945847
  Influence
                οf
                        purinoceptor
                                         antagonism
                                                         on
pentaphosphate-induced hypotension in anesthetized rats.
  Steinmetz M; van Le T; Hollah P; Gabriels G; Hohage H; Rahn K H;
Schlatter E
  Medizinische
                Poliklinik,
                              Experimentelle
                                               Nephrologie,
Wilhelms-Universitat, Munster, Germany.
  Journal of pharmacology and experimental therapeutics (UNITED STATES)
Sep 2000, 294 (3) p963-8, ISSN 0022-3565 Journal Code: 0376362
 Document type: Journal Article
 Languages: ENGLISH
 Main Citation Owner: NLM
 Record type: Completed
  ... A(2) purinoceptor antagonist 3, 7-dimethyl-1-propargylxanthine. The
hypertensive effect by the prototype P2X
                                                      receptor
alphabeta-methylene ATP was inhibited by pyridoxal-phosphate-6-azophenyl
-2',4'-disulfonic acid, too. Purinoceptor antagonists reduced the maximal
? ds
Set
       Items
               Description
S1
           Ω
               P2X (2) RECEPTOR
S2
          10
               P()2()X (N2) RECEPTOR
        5539
               P2X (N2) RECEPTOR
        5539
               "P2X" (N2) RECEPTOR
               P2X4 (N2) RECEPTOR
         277
         383
               S3 (N2) ANTAGONIST
         383
               S4 (N2) ANTAGONIST
         390
               S3 (N2) AGONIST
         390
               S4 (N2) AGONIST
           2
               S5 (N2) ANTAGONIST
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diadenosine

Westfalische

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S3
S4
S5 -
S6
S7
S8
S9
S10
S11
            15
                 S5 (N2) AGONIST
                 S10 (N5) INHIBIT?
S12
            1
S13
             0
                 S11 (N5) INHIBIT?
        19158
S14
                 NEUROPATHIC (N2) PAIN
                 TACTILE (N2) ALLODYNIA
S15
         1570
S16
          932
                 S14 AND S15
          344
S17
                 RD (unique items)
                 S17 AND S5
S18
            2
S19
          148
                 17 AND S4
S20
            76
                 RD (unique items)
S21
             5
                 S17 AND S4
S22
             5
                 RD (unique items)
S23
             0
                 S5 AND ION()FLUX
S24
             9
                 S4 AND ION()FLUX
S25
             5
                 RD (unique items)
S26
          194
                 S9 AND INHIBIT?
S27
                 S9 (N5) INHIBIT?
? s s7(N5)inhibit?
Processed 20 of 29 files ...
Processing
Completed processing all files
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383 S7

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6944746 INHIBIT?
27 S7(N5)INHIBIT?
     S28
...completed examining records
             10 RD (unique items)
? t/3, k/all
>>>KWIC option is not available in file(s): 399
29/3,K/1 (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
0014527335 BIOSIS NO.: 200300496054
Unusual absence of endothelium-dependent or -independent vasodilatation to
  purines or pyrimidines in the rat renal artery.
AUTHOR: Knight Gillian E; Oliver-Redgate Rachel; Burnstock Geoffrey
  (Reprint)
AUTHOR ADDRESS: Medical School, Autonomic Neuroscience Institute, Royal
  Free and University College, Rowland Hill Street, Royal Free Campus,
  London, NW3 2PF, UK**UK
AUTHOR E-MAIL ADDRESS: g.burnstock@ucl.ac.uk
JOURNAL: Kidney International 64 (4): p1389-1397 October 2003 2003
MEDIUM: print
ISSN: 0085-2538 (ISSN print)
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
... ABSTRACT: of P2X and P2Y receptor subtypes was performed. Results:
 Electrical field stimulation induced vasoconstriction, partially
  inhibited by the P2X
                         receptor antagonist,
 pyridoxalphosphate-6-azophenyl-2',4'-disulfonic acid, and predominantly
 by prazosin. Exogenous NA and ATP...
 29/3,K/2
             (Item 2 from file: 5)
DIALOG(R)File, 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
0013376178 BIOSIS NO.: 200100548017
Purinergic co-transmission in postnatal mouse lateral hypothalamus in vitro
AUTHOR: Jo Y H (Reprint); Role L W (Reprint)
AUTHOR ADDRESS: Dept Anat and Cell Biol Ctr Neruobiol, Columbia Univ Col
  Physicians and Surgeons, New York, NY, USA**USA
JOURNAL: Society for Neuroscience Abstracts 27 (2): p1572 2001 2001
MEDIUM: print
CONFERENCE/MEETING: 31st Annual Meeting of the Society for Neuroscience
San Diego, California, USA November 10-15, 2001; 20011110
ISSN: 0190-5295
DOCUMENT TYPE: Meeting; Meeting Abstract
RECORD TYPE: Abstract
LANGUAGE: English
DESCRIPTORS:
 CHEMICALS & BIOCHEMICALS: ... P2X receptor antagonist , current
    inhibitor; ...
... P2X receptor antagonist, current inhibitor
```

29/3,K/3 (Item 3 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2004 BIOSIS. All rts. reserv.

0012536936 BIOSIS NO.: 200000255249

Effect of age on the responses of rat bladder detrusor strips to adenosine triphosphate

AUTHOR: Kageyama S (Reprint); Fujita K; Suzuki K; Shinbo H; Masuda N;

AUTHOR ADDRESS: 3600 Handa-cho, Hamamatsu-shi, Shizuoka-ken, Japan**Japan

JOURNAL: BJU International 85 (7): p899-904 May, 2000 2000

MEDIUM: print ISSN: 1464-4096

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: did not significantly inhibit the phasic contraction, but reduced the postwashout contraction. PPADS (a selective P2X receptor antagonist) did not inhibit either contraction. Indomethacin (a prostaglandin synthesis inhibitor) had no effect on the phasic contraction but...

29/3,K/4 (Item 4 from file: 5)

DIALOG(R) File 5:Biosis Previews(R) (c) 2004 BIOSIS. All rts. reserv.

0011806702 BIOSIS NO.: 199900066362

ATP P2X receptors and sensory synaptic transmission between primary afferent fibers and spinal dorsal horn neurons in rats

AUTHOR: Li Ping; Calejesan Amelita A; Zhuo Min

AUTHOR ADDRESS: Dep. Ansthesiol., Washington Univ., Sch. Med., St. Louis, MO 63110-1093, USA**USA

JOURNAL: Journal of Neurophysiology (Bethesda) 80 (6): p3356-3360 Dec., 1998 1998

MEDIUM: print ISSN: 0022-3077

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: the lumbar spinal cord.

Pyridoxal-phosphate-6-azophenyl-2',4'-disulfonic acid (PPADS), a selective P2X receptor antagonist, produced an inhibitory modulatory effect on fast EPSCs and altered responses to paired-pulse stimulation, suggesting the involvement...

29/3,K/5 (Item 5 from file: 5)

DIALOG(R) File 5: Biosis Previews(R) (c) 2004 BIOSIS. All rts. reserv.

(0) 2001 B10010. H11 100. 100011.

0010688008 BIOSIS NO.: 199799322068

P2X purinoceptors in cultured myenteric neurons of guinea-pig small intestine

AUTHOR: Zhou Xiaoping; Galligan James J (Reprint)

AUTHOR ADDRESS: Dep. Pharmacology Toxicology, Michigan State Univ., East

Lansing, MI, USA**USA

JOURNAL: Journal of Physiology (Cambridge) 496 (3): p719-729 1996 1996

ISSN: 0022-3751

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

...ABSTRACT: 100 mu-M) partly inhibited f-EPSCs in 28% of neurons. Hexamethonium-resistant fEPSCs were inhibited by 97 +- 2% by the P2X receptor antagonist, pyridoxal-phosphate-6-azophenyl-2',4'-disulphonic acid (PPADS, 10 mu-M). 2. ATP caused...

29/3,K/6 (Item 1 from file: 34)

DIALOG(R) File 34:SciSearch(R) Cited Ref Sci (c) 2004 Inst for Sci Info. All rts. reserv.

12511896 Genuine Article#: 774FM No. References: 31

Title: Desensitization masks nanomolar potency of ATP for the P2X(1) receptor

Author(s): Rettinger J; Schmalzing G (REPRINT)

Corporate Source: Rhein Westfal TH Aachen, Sch Med, Dept Mol

Pharmacol, Wendlingweg 2/D-52074 Aachen//Germany/ (REPRINT); Rhein

Westfal TH Aachen, Sch Med, Dept Mol Pharmacol, D-52074 Aachen//Germany/;

Max Planck Inst Biophys, D-60439 Frankfurt//Germany/

Journal: JOURNAL OF BIOLOGICAL CHEMISTRY, 2004, V279, N8 (FEB 20), P 6426-6433

ISSN: 0021-9258 Publication date: 20040220

Publisher: AMER SOC BIOCHEMISTRY MOLECULAR BIOLOGY INC, 9650 ROCKVILLE

PIKE, BETHESDA, MD 20814-3996 USA

Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

...Abstract: P2X(2)/P2X(1) chimera and the P2X(1) receptor possess virtually identical sensitivity to inhibition by the P2X (1) receptor -selective antagonist NF279, a suramin analog. These results suggest that the P2X(1), ectodomain confers nanomolar ATP...

29/3,K/7 (Item 2 from file: 34)

DIALOG(R) File 34:SciSearch(R) Cited Ref Sci (c) 2004 Inst for Sci Info. All rts. reserv.

11302902 Genuine Article#: 635LK No. References: 58

Title: Mechanisms of P2X(7) receptor-mediated ERK1/2 phosphorylation in human astrocytoma cells

Author(s): Gendron FP (REPRINT); Neary JT; Theiss PM; Sun GY; Gonzalez FA; Weisman GA

Corporate Source: Univ Missouri, Dept Biochem, M121 Med Sci

Bldg/Columbia//MO/65212 (REPRINT); Univ Missouri, Dept

Biochem, Columbia//MO/65212; Univ Miami, Vet Affairs Med Ctr, Res Serv,

Sch Med, Miami//FL/33125; Univ Miami, Dept Pathol, Sch

Med, Miami//FL/33125; Univ Miami, Dept Biochem & Mol Biol, Sch

Med, Miami//FL/33125; Univ Miami, Neurosci Program, Sch

Med, Miami//FL/33125; Univ Puerto Rico, Dept Chem, Rio Piedras//PR/00931

Journal: AMERICAN JOURNAL OF PHYSIOLOGY-CELL PHYSIOLOGY, 2003, V284, N2 (FEB), PC571-C581

ISSN: 0363-6143 Publication date: 20030200

Publisher: AMER PHYSIOLOGICAL SOC, 9650 ROCKVILLE PIKE, BETHESDA, MD 20814

USA

Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

... Abstract: cells overexpressing the recombinant rat P2X(7) receptor

(rP2X(7)-R), a response that was inhibited by the P2X (7) receptor
antagonist , oxidized ATP. Other results suggest that
rP2X(7)-R-mediated ERK1/2 phosphorylation was linked...

29/3,K/8 (Item 3 from file: 34)
DIALOG(R)File 34:SciSearch(R) Cited Ref Sci
(c) 2004 Inst for Sci Info. All rts. reserv.

10093469 Genuine Article#: 483EP No. References: 22

Title: Differential responses to ATP gamma S in the mesenteric and hindlimb vascular bed of the cat

Author(s): Shah MK (REPRINT); Champion HC; Bivalacqua TJ; Kadowitz PJ Corporate Source: 10 Summer St, Apt 303S/Malden//MA/02148 (REPRINT); Tulane Univ, Sch Med, Dept Pharmacol, New Orleans//LA/70112

Journal: LIFE SCIENCES, 2001, V69, N21 (OCT 12), P2561-2571

ISSN: 0024-3205 Publication date: 20011012

Publisher: PERGAMON-ELSEVIER SCIENCE LTD, THE BOULEVARD, LANGFORD LANE, KIDLINGTON, OXFORD OX5 1GB, ENGLAND

Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

... Abstract: In the mesenteric vascular bed the pressor response to ATP gammaS was blocked by a P2X (1) receptor antagonist. Also an inhibitor of nitric oxide synthase enhanced the vasoconstrictive responses to ATP gammaS. However, the vasodepressor response...

29/3,K/9 (Item 4 from file: 34)
DIALOG(R)File 34:SciSearch(R) Cited Ref Sci
(c) 2004 Inst for Sci Info. All rts. reserv.

06939195 Genuine Article#: 105WG No. References: 17
Title: 2',3'-O-(2,4,6-trinitrophenyl)adenosine 5'-triphosphate (TNP-ATP) a nanomolar affinity antagonist at rat mesenteric artery P2X receptor
ion channels

Author(s): Lewis CJ; Surprenant A; Evans RJ (REPRINT)

Corporate Source: UNIV LEICESTER, DEPT CELL PHYSIOL & PHARMACOL, MED SCI BLDG, UNIV RD/LEICESTER LE1 9HN/LEICS/ENGLAND/ (REPRINT); UNIV LEICESTER, DEPT CELL PHYSIOL & PHARMACOL/LEICESTER LE1 9HN/LEICS/ENGLAND/; GENEVA BIOMED RES INST,/GENEVA//SWITZERLAND/

Journal: BRITISH JOURNAL OF PHARMACOLOGY, 1998, V124, N7 (AUG), P1463-1466 ISSN: 0007-1188 Publication date: 19980800

Publisher: STOCKTON PRESS, HOUNDMILLS, BASINGSTOKE RG21 6XS, HAMPSHIRE, ENGLAND

Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

... Abstract: smooth muscle cells and contractions of whole artery rings.

2 The selective P2X(1) and **P2X** (3) **receptor antagonist** TNP-ATP **inhibited** P2X receptor mediated inward currents in response to 3 mu M alpha, beta-meATP (an...

29/3,K/10 (Item 1 from file: 144) DIALOG(R)File 144:Pascal (c) 2004 INIST/CNRS. All rts. reserv.

16063767 PASCAL No.: 03-0213415

Mechanisms of P2X SUB 7 receptor-mediated ERK1/2 phosphorylation in human astrocytoma cells

GENDRON Fernand-Pierre; NEARY Joseph T; THEISS Patty M; SUN Grace Y; GONZALEZ Fernando A; WEISMAN Gary A

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... the recombinant rat P2X SUB 7 receptor (rP2X SUB 7 -R), a response that was inhibited by the P2X SUB 7 receptor antagonist, oxidized ATP. Other results suggest that rP2X SUB 7 -R-mediated ERK1/2 phosphorylation was...?

```
Set
        Items
                 Description
            0
                 P2X(2) RECEPTOR
S1
S2
           10
                 P()2()X (N2) RECEPTOR
                 P2X (N2) RECEPTOR
         5539
S3
         5539
                 "P2X" (N2) RECEPTOR
S4
          277
                 P2X4 (N2) RECEPTOR
$5
          383
                 S3 (N2) ANTAGONIST
S6
          383
                 S4 (N2) ANTAGONIST
s7
          390
                 S3 (N2) AGONIST
S8
                 S4(N2)AGONIST
S9
          390
                 S5 (N2) ANTAGONIST
S10
            2
           15
                 S5 (N2)AGONIST
S11
                 S10(N5)INHIBIT?
S12
            1
             0
                 S11(N5)INHIBIT?
S13
        19158
                 NEUROPATHIC (N2) PAIN
S14
S15
         1570
                 TACTILE (N2) ALLODYNIA
                 S14 AND S15
S16
           932
                 RD (unique items)
S17
          344
             2
                 S17 AND S5
S18
                 17 AND S4
S19
           148
                 RD (unique items)
            76
S20
                 S17 AND S4
             5
S21
             5
                 RD (unique items)
S22
             0
                 S5 AND ION()FLUX
S23
             9
S24
                 S4 AND ION()FLUX
             5
                 RD (unique items)
S25
           194
                 S9 AND INHIBIT?
S26
                 S9 (N5) INHIBIT?
             5
S27
            27
                 S7 (N5) INHIBIT?
S28
S29
            10
                 RD (unique items)
? s s14(s)s7
            19158
                   S14
                   S7
              383
     S30
                0
                   S14(S)S7
? s s14(N10)s7
            19158
                   S14
              383
                   S7
     S31
                0
                   S14(N10)S7
? s s14(n10)s8
            19158
                   S14
              390
                   S8
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? ds
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Items
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Set
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S1
                P()2()X (N2) RECEPTOR
           10
S2
                P2X (N2) RECEPTOR
S3
         5539
S4
         5539
                "P2X" (N2) RECEPTOR
                P2X4 (N2) RECEPTOR
S5
          277
S6
          383
                S3 (N2) ANTAGONIST
          383
                S4 (N2) ANTAGONIST
S7
          390
                S3 (N2) AGONIST
S8
          390
                S4(N2)AGONIST
S9
                S5 (N2) ANTAGONIST
S10
            2
                S5 (N2) AGONIST
S11
           15
                S10 (N5) INHIBIT?
S12
            1
                S11 (N5) INHIBIT?
S13
            0
S14
        19158
                NEUROPATHIC (N2) PAIN
         1570
                TACTILE (N2) ALLODYNIA
S15
          932
                S14 AND S15
S16
                RD (unique items)
          344
S17
                S17 AND S5
            2
S18
                17 AND S4
S19
          148
           76
                RD (unique items)
S20
                S17 AND S4
            5
S21
                RD (unique items)
S22
                S5 AND ION()FLUX
S23
            0
                S4 AND ION() FLUX
S24
S25
            5
                RD (unique items)
          194
                 S9 AND INHIBIT?
S26
            5
                 S9 (N5) INHIBIT?
S27
           27
                 S7 (N5) INHIBIT?
S28
                 RD (unique items)
           10
S29
S30
            0
                 S14(S)S7
            0
                 S14(N10)S7
S31
S32
            0
                 S14(N10)S8
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     $1.35 Estimated cost File6
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            $42.33
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        $0.31
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 $1.10
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